Patent Attorney Docket No. 74239

## **AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions and listings of claims in the application:

## **LISTING OF CLAIMS:**

- 1. (Currently amended): A method of producing a molecularly-imprinted material, comprising:
  - (a) synthesizing a peptide, eligosaccharide or eligonucleotide on a
    disposable surface modified support to produce a support surfaceattached peptide, eligosaccharide or eligonucleotide;
  - (b) providing a selected monomer mixture;
  - (c) contacting said monomer mixture with said support surface-attached peptide, oligosaccharide er oligonuelectide;
  - (d) initiating polymerisation or at least one crosslinking reaction;
  - (e) dissolving or degrading said support surface-attached peptide,
     oligosaccharide or oligonucleotide and said support; and
  - (f) obtaining said molecularly imprinted material.
- 2. (Original): A method according to claim 1, wherein said peptide of step (c) is a peptide epitope.
- 3. (Original): A method according to claim 1, wherein step (f) is conducted with the aid of at least one factor consisting of crosslinking agents, heat, and ultraviolet irradiation.
- 4. (Original): A method according to claim 1, wherein said peptide is selected from the group consisting of FMOC-Phe-Gly-Si, H-Phe-Gly-Si, FMOC-Phe-Si, BOC-Gly-Si, H-Gly-Si, FMOC-Phe-Gly-OH, FMOC-Phe-OH, BOC-Phe-OH, H-Phe-PNA, H-Phe-O-Me, H-Phe-OtBu, BOC-Gly-OH, H-Phe-Gly-NH<sub>2</sub>, H-Phe-Gly-Gly-Phe-OH, FMOC-Phe-OH, H-Gly-Phe-Oll, and Nociceptin.

Patent Attorney Docket No. 74239

- 5. (Currently amended): A method according to claim 1, wherein said disposable surface activated modified support is a silane-modified silica or controlled pore glass (CPG).
- 6. (Original): A method according to claim 1, wherein said monomer mixture comprises monomers selected from the group consisting of styrene/divinyl benzene, methacrylates, acrylates, acrylamides, methacrylamides and combinations thereof.
- 7. (Withdrawn): A method of using a molecularly-imprinted material, comprising:

producing a molecularly-imprinted material according to claim 1; and using said molecularly-imprinted material as an affinity phase for the separation of biological macromolecules or oligomers.

- 8. (Withdrawn): A method according to claim 7, wherein said biological macromolecules or oligomers are selected from the group consisting of peptides, polypeptides, oligopeptides, proteins, nucleic acids, oligonucleotides, polynucleotides, saccharides, oligosaccharides, and polysaccharides.
- 9. (Withdrawn): A chromatographic stationary phase, comprising a molecularly imprinted material produced according to claim 1, wherein said peptide, oligosaccharide or oligonucleotide of step (c) is selected from the group consisting of FMOC-Phe-Gly-Si, H-Phe-Gly-Si, FMOC-Phe-Si, BOC-Gly-Si, H-Gly-Si, FMOC-Phe-Gly-OH, FMOC-Phe-OH, BOC-Phe-OH, H-Phe-PNA, H-Phe-O-Me, H-Phe-OtBu, BOC-Gly-OH, H-Phe-Gly-NH<sub>2</sub>, H-Phe-Gly-Gly-Phe-OH, FMOC-Phe-OH, and H-Gly-Phe-OH, and Nociceptin.